## **Amendments**

## Amendments to the Claims

Please amend the claims as shown below in the List of Claims

## List of Claims

- 1-12. (Canceled).
- 13. (Currently Amended) A process for the production of an L-amino acid chosen from the group consisting of L-threonine, L-isoleucine, L-valine, L-methionine, L-homoserine and L-lysine comprising:
  - a) fermenting a bacterium comprising an overexpressed endogenous DNA sequence encoding the galactose-proton symporter protein in said bacterium, in a fermentation medium under conditions suitable for the production of said L-amino acid, wherein:
    - i) said bacterium is of an Enterobacteriaceae family;
    - ii) said galactose-proton symporter protein comprises the amino acid sequence of SEQ ID NO:4 and is encoded by the nucleotide sequence of residues 33-1427 of SEQ ID NO:3;
    - iii) said L-amino acid is produced from glucose, saccharose, lactose, fructose, molasses, starch, cellulose or from glycerine and ethanol;
    - iv) said overexpression is achieved by increasing the copy number of said DNA or by operably linking said DNA to a promoter changing a promoter normally found in a galP gene; and
  - b) allowing said L-amino acid to become enriched in said bacteria or said fermentation medium.
- 14. (Previously Presented) The process of claim 13, wherein said galactose-proton symporter protein consists of the amino acid sequence of SEQ ID NO:4.

- 15. (Currently Amended) The process of claim 14, wherein said DNA sequence encoding the galactose-proton symporter protein consists of the nucleotide sequence of residues 33-1427 of SEQ ID NO:3.
- 16. (Currently Amended) The process of claim 13, wherein said DNA sequence encoding the galactose-proton symporter protein consists of the nucleotide sequence of residues 33-1427 of SEQ ID NO:3.
- 17. (Previously Presented) The process of claim 13, wherein overexpression is achieved by increasing the copy number of said DNA.
- 18. (Previously Presented) The process of claim 13, wherein said L-amino acid is L-threonine.

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- 19. (Previously Presented) The process of any one of claims 13-16, further comprising isolating said L-amino acid along with some or all of the constituents of said fermentation medium and/or the biomass in said fermentation medium.
- 20. (Previously presented) The process of claim 19, wherein said L-amino acid is L-threonine.
- 21. (Currently Amended) The process of claim 13, wherein said microorganism overexpresses one or more genes selected from the group consisting of:
  - a) the <u>a</u> thrABC operon coding for aspartate kinase, homoserine dehydrogenase, homoserine kinase and threonine synthase;
  - b) the a pyc gene coding for pyruvate carboxylase;
  - c) the a pps gene coding for phosphoenolpyruvate synthase;
  - d) the a ppc gene coding for phosphoenolpyruvate carboxylase;

- e) the pntA and pntB genes coding for transhydrogenase,
- f) the a rhtB gene which imparts homoserine resistance;
- g) the a mgo gene coding for malate:quinone oxidoreductase;
- h) the a rhtC gene which imparts threonine resistance;
- i) the thrE gene coding for threonine export protein;
- j) the a gdhA gene coding for glutamate dehydrogenase;
- k) the a glk gene coding for glucokinase;
- 1) the a hns gene coding for DNA binding protein HLP-II;
- m) the a pgm gene coding for phosphoglucomutase;
- n) the <u>a</u> fba gene coding for fructose biphosphate aldolase;
- o) the <u>a</u> ptsH gene coding for phosphohistidine protein hexose phosphotransferase;
- p) the a ptsI gene coding for enzyme I in the phosphotransferase system;
- q) the <u>a</u> crr gene coding for the glucose-specific IIA component;
- r) the a ptsG gene coding for the glucose-specific IIBC component;
- s) the <u>a</u> lrp gene coding for a regulator in the leucine regulon;
- t) the a csrA gene coding for the global regulator Csr;
- u) the <u>a</u> fadR gene coding for a regulator in the fad regulon;
- v) the a iclR gene coding for a regulator in central intermediary metabolism;
- w) the a mopB gene coding for the 10 KDa chaperone;
- x) the <u>a</u> ahpC gene coding for the small sub-unit of alkyl hydroperoxide reductase;
- y) the a ahpF gene coding for the large sub-unit of alkyl hydroperoxide reductase;
- z) the a cysK gene coding for cysteine synthase A;
- aa) the a cysB gene coding for the regulator in the cys regulon;
- bb) the a cysJ gene coding for the flavoprotein in NADPH sulfite reductase;
- cc) the a cysI gene coding for haemoprotein in NADPH sulfite reductase;
- dd) the a cysH gene coding for adenylylsulfate reductase;
- ee) the a phoB gene coding for the positive regulator PhoB in the pho regulon;

- ff) the a phoR gene coding for the sensor protein in the pho regulon;
- gg) the a phoE gene coding for protein E in the outer cell membrane;
- hh) the a pykF gene coding for the pyruvate kinase I stimulated by fructose;
- ii) the a pfkB gene coding for 6-phosphofructokinase II;
- jj) the a malE gene coding for periplasmatic binding protein in maltose transport;
- kk) the a sodA gene coding for superoxidedismutase;
- 11) the a rseA gene coding for a membrane protein with anti-sigmaE activity;
- mm) the a rseC gene coding for a global regulator in the sigmaE factor;
- nn) the a sucA gene coding for the decarboxylase sub-unit of 2-ketoglutarate dehydrogenase;
- oo) the a sucB gene coding for the dihydrolipoyl-transsuccinase E2 subunit of 2-ketoglutarate dehydrogenase;
- pp) the <u>a</u> sucC gene coding for the  $\beta$ -subunit of succinyl-CoA synthetase;
- qq) the <u>a</u> sucD gene coding for the  $\alpha$ -subunit in succinyl-CoA synthetase;
- rr) the a adk gene coding for adenylate kinase;
- ss) the <u>a</u> hdeA gene coding for a periplasmatic protein with a chaperonin-like function;
- tt) the a hdeB gene coding for a periplasmatic protein with a chaperonin-like function;
- uu) the a icd gene coding for isocitrate dehydrogenase;
- vv) the a mglB gene coding for periplasmatic, galactose-binding transport protein;
- ww) the a lpd gene coding for dihydrolipoamide dehydrogenase;
- xx) the <u>a</u> aceE gene coding for the E1 component of pyruvate dehydrogenase complex;
- yy) the <u>a</u> aceF gene coding for the E2 component of pyruvate dehydrogenase complex;
- zz) the a pepB gene coding for aminopeptidase B;
- aaa) the a aldH gene coding for aldehyde dehydrogenase;
- bbb) the a bfr gene coding for the iron storage homoprotein;

- ccc) the a udp gene coding for uridine phosphorylase; and
  ddd) the a rseB gene coding for the regulator of sigmaE factor activity;
  wherein said overexpression is achieved by one or more methods selected from the
  group consisting of increasing copy number, using a strong promoter, and mutating the
  ribosome binding site.
- 22. (Currently Amended) The process of claim 13, wherein at least one gene in said microorganism is attenuated deleted, said gene being selected from the group consisting of:
  - a) the <u>a</u> tdh gene coding for threonine dehydrogenase;
  - b) the a mdh gene coding for malate dehydrogenase;
  - c) the a gene product of the open reading frame (ORF) yifA;
  - d) the a gene product of the open reading frame (ORF) ytfP;
  - e) the a pckA gene coding for the enzyme phosphoenol-pyruvate carboxykinase;
  - f) the a poxB gene coding for pyruvate oxidase;
  - g) the <u>a</u> aceA gene coding for isocitrate lyase;
  - h) the <u>a</u> dgsA gene coding for the DgsA regulator in the phosphotransferase system;
  - $\frac{1}{h}$  the <u>a</u> fruR gene coding for fructose repressor;
  - <u>ji</u>) the a rpoS gene coding for the sigma<sup>38</sup>-Factor;
  - ki) the a aspA gene coding for aspartate ammonium lyase; and
  - $\frac{1}{k}$  the <u>a</u> aceB gene coding for malate synthase A gene.
- 23. (Currently amended) A process for the production of an L-amino acid chosen from the group consisting of L-threonine, L-isoleucine, L-valine, L-methionine, L-homoserine and L-lysine comprising:
  - a) fermenting a bacterium comprising an overexpressed endogenous DNA sequence encoding the galactose-proton symporter protein in said bacterium,

in a fermentation medium under conditions suitable for the production of said L-amino acid, wherein:

- i) said bacterium is of an Enterobacteriaceae family and transports

  glucose by a PEP-dependent phosphotransferase (PTS) pathway

  comprises PTS enzymes;
- ii) said galactose-proton symporter protein comprises the amino acid sequence of SEQ ID NO:4;
- said L-amino acid is produced from glucose, saccharose, lactose,fructose, molasses, starch, cellulose or from glycerine and ethanol;
- iv) said overexpression is achieved by increasing the copy number of said DNA or by operably linking said DNA to a promoter changing a promoter normally found in a galP gene; and
- b) allowing said L-amino acid to become enriched in said bacteria or said fermentation medium.
- 24. (Previously presented) The process of claim 23, further comprising isolating said L-amino acid along with some or all of the constituents of said fermentation medium and/or the biomass in said fermentation medium.
- 25. (Previously Presented) The process of claim 24, wherein said bacterium is selected from the group consisting of: Escherichia coli H4581; Escherichia coli VNIIgenetika MG442; Escherichia coli VNIIgenetika M1; Escherichia coli VNIIgenetika 472T23; Escherichia coli BKIIM B-3996; Escherichia coli kat 13; and Escherichia coli KCCM-10132.
- 26. (Previously presented) The process of claim 25, wherein said L-amino acid is L-threonine.